

-continued

Ser	Ser	Ser	Leu	Gly	Thr	Gln	Thr	Tyr	Ile	Cys	Asn	Val	Asn	His	Lys
		195					200					205			
Pro	Ser	Asn	Thr	Lys	Val	Asp	Lys	Lys	Val	Glu	Pro	Lys	Ser	Cys	Asp
	210					215					220				
Lys	Thr	His	Thr	Cys	Pro	Pro	Cys	Pro	Ala	Pro	Glu	Leu	Leu	Gly	Gly
	225				230					235					240
Pro	Ser	Val	Phe	Leu	Phe	Pro	Pro	Lys	Pro	Lys	Asp	Thr	Leu	Met	Ile
			245						250					255	
Ser	Arg	Thr	Pro	Glu	Val	Thr	Cys	Val	Val	Val	Asp	Val	Ser	His	Glu
			260					265					270		
Asp	Pro	Glu	Val	Lys	Phe	Asn	Trp	Tyr	Val	Asp	Gly	Val	Glu	Val	His
		275					280					285			
Asn	Ala	Lys	Thr	Lys	Pro	Arg	Glu	Glu	Gln	Tyr	Asn	Ser	Thr	Tyr	Arg
	290					295					300				
Val	Val	Ser	Val	Leu	Thr	Val	Leu	His	Gln	Asp	Trp	Leu	Asn	Gly	Lys
	305				310					315					320
Glu	Tyr	Lys	Cys	Lys	Val	Ser	Asn	Lys	Ala	Leu	Pro	Ala	Pro	Ile	Glu
			325						330					335	
Lys	Thr	Ile	Ser	Lys	Ala	Lys	Gly	Gln	Pro	Arg	Glu	Pro	Gln	Val	Tyr
		340						345					350		
Thr	Leu	Pro	Pro	Ser	Arg	Glu	Glu	Met	Thr	Lys	Asn	Gln	Val	Ser	Leu
		355					360					365			
Thr	Cys	Leu	Val	Lys	Gly	Phe	Tyr	Pro	Ser	Asp	Ile	Ala	Val	Glu	Trp
	370					375					380				
Glu	Ser	Asn	Gly	Gln	Pro	Glu	Asn	Asn	Tyr	Lys	Thr	Thr	Pro	Pro	Val
	385				390					395					400
Leu	Asp	Ser	Asp	Gly	Ser	Phe	Phe	Leu	Tyr	Ser	Lys	Leu	Thr	Val	Asp
			405					410					415		
Lys	Ser	Arg	Trp	Gln	Gln	Gly	Asn	Val	Phe	Ser	Cys	Ser	Val	Met	His
		420						425					430		
Glu	Ala	Leu	His	Asn	His	Tyr	Thr	Gln	Lys	Ser	Leu	Ser	Leu	Ser	Pro
		435					440					445			

Gly

What is claimed is:

1. A method for purifying a polypeptide from a composition comprising the polypeptide and a contaminant, which method comprises the following steps performed sequentially:

- (a) loading the composition onto an ion exchange material;
- (b) washing the ion exchange material with wash buffer using a multi-slope gradient, wherein the multi-slope gradient comprises two or more segments of linear salt gradients with different slopes wherein the slope is greater in the first segment than in any additional segments, the increase in the salt concentration of the wash buffer is greater in the first segment of the multi-slope gradient wash than in subsequent segments, and each segment of the multi-slope wash ends when a predetermined polypeptide concentration is measured in the flowthrough, wherein each segment of the multi-slope gradient has a progressively shallower slope; and
- (c) eluting the polypeptide from the ion exchange material.

2. The method of claim 1, additionally comprising the step between steps (b) and (c) of washing the column with from 0.4 to 1 column volumes of wash buffer.

3. The method of claim 2 wherein the wash buffer has the composition of the wash buffer at the end of step (b).

4. The method of claim 1, 2, or 3, further comprising subjecting the composition comprising the polypeptide to one or more further purification steps so as to obtain a homogenous preparation of the polypeptide.

5. The method of claim 1, wherein the polypeptide is an antibody.

6. The method of claim 5, wherein the antibody is an anti-HER2 antibody.

7. The method of claim 6, wherein the anti-HER2 antibody comprises the light chain amino acid sequence of SEQ ID NO: 1 and the heavy chain amino acid sequence of SEQ ID NO: 2.

8. The method of claim 7, wherein the antibody is rhuMAb HER2 comprising the light chain amino acid sequence of SEQ ID NO: 1, and the heavy chain amino acid sequence of SEQ ID NO: 2, and wherein the contaminants comprise a deamidated variant having Asn30 in CDR1 of either or both of the light chain variable regions (VL) converted to aspartate.

9. The method of claim 7 or 8, wherein the multi-slope gradient comprises three segments.